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SUPPLEMENTAL RESPONSE TO FINAL OFFICE ACTION

admixing or binding a biologically-active factor selected from the group 1) consisting of TNF-\alpha and lymphotoxin and a target molecule with a colloidal metal to form a composition; and

2) administering the composition to the human or animal.

The method of Claim 8, wherein the biologically-active (Twice Amended) factor is tumor necrosis factor.

(Thrice Amended) A method of treating a human or animal with a cancer or immune disease comprising administering to the human or animal with the cancer or immune disease a therapeutically effective amount of a composition capable of targeting a particular tissue comprising a biologically-active factor selected from the group consisting of TNF-\alpha and lympotoxin and a target molecule admixed with or bound to a colloidal metal.

The method of Claim 18, wherein the biologically-active ("INFA") 16. (Twice Amended) factor is tumor necrosis facto

1/9.17 (Twice Amended) A method for the delivery of more than one biologicallyactive factor comprising administering to a human or animal a composition comprising more than one biologically-active factor selected from the group consisting of TNF-a and lymphotoxin and a target molecule admixed with or bound to a colloidal metal.

(Twice Amended) The method of Claim 19 wherein the biologically active factor is tumor necrosis factor ("TNFa").

(Twice Amended) A method for the targeted delivery of one or more 21. biologically-active factors, comprising administering to a human or animal a composition comprising two or more biologically-active factors selected from the group consisting of TNF-\alpha and lymphotoxin admixed with or bound to colloidal metal wherein at least one of the biologically-active factors is a target molecule

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capable of binding a receptor on a dell membrane and wherein at least one of the biologically-active factors is released from the composition in vivo.

(Twice Amended) The method of Claim 21 wherein the biologically-active factor is tumor necrosis factor ("TNF α ").

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- (Twice Amended) A method of treating a human or animal with cancer or an immune disease comprising administering to the human or animal a composition comprising two or more biologically-active factor selected from the group consisting of TNF-or and lymphotoxin admixed with or bound to a colloidal metal, wherein at least one of the biologically-active factors is a target molecule capable of binding a receptor on a cell membrane.
- 25. (Twice Amended) The method of Claim 24 wherein the biologically-active factor is tumor necrosis factor ("TNFα").

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- (Amended) The method of Claim 21, wherein the composition further comprises additional biologically active factors admixed with or bound to the colloidal metal.
- 33. (Amended) A method of treating a human or animal with a cancer comprising administering to the human or animal with the cancer a therapeutically effective amount of a composition comprising a biologically-active factor selected from the group consisting of TNF-α and lymphotoxin admixed with or bound to a colloidal metal.

34. (Amended) A method of treating a human or animal with a cancer or immune disease comprising administering to the human or animal with the cancer or immune disease a therapeutically effective amount of a composition comprising a biologically-active factor which is tumor necrosis factor ("TNFα") admixed with or bound to a colloidal metal.

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